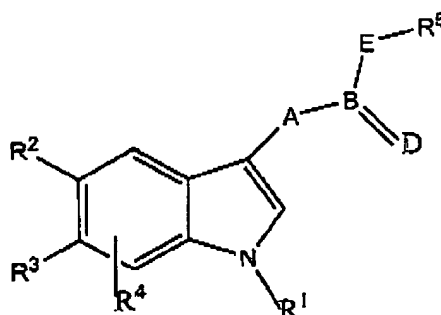


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IN THE CLAIMSCLAIMS

1. A method for the treatment of a skin disease comprising topically administering a subject in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof:



1

in which

R¹ is

(i) -C₁₋₁₂-alkyl, straight-chain or branched-chain or -C₂-C₁₂ alkenyl, mono- or polyunsaturated,

optionally mono- or polysubstituted by -OH, -SH, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NHC₆₋₁₄ aryl, -N(C₆₋₁₄aryl)₂, -N(C₁₋₁₄aryl)₂, -N(C₁₋₆alkyl)(C₆₋₁₄aryl), -NHCOR⁶, -NO₂, -CN, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -O-C₆₋₁₄-aryl, -O(CO)R⁶, -S-C₁₋₆-alkyl, -S-C₆₋₁₄aryl, -SOR⁶, -SO₃H, -SO₂R⁶, -OSO₂C₁₋₆-alkyl, -OSO₂C₆₋₁₄aryl, -(CS)R⁶, -COOH, -(CO)R⁶, mono-, bi or tricyclic saturated or mono- or polyunsaturated carbocycles having 3-14 ring members, mono-, bi- or

tricyclic saturated or mono- or polyunsaturated heterocycles having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S, wherein the C₆₋₁₄aryl groups and the carbocyclic and heterocyclic substituents for their part can optionally be mono- or polysubstituted by R⁴,

(ii) a mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycle having 3-14 ring members or a mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycle having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S, or a carbo- or heterocyclic saturated or mono- or polyunsaturated spirocycle having 3-10 ring members, where heterocyclic systems contain 1-6 heteroatoms, which are preferably N, O and S, optionally mono- or polysubstituted by -OH, -SH, -NH₂, NHC₁₋₆ alkyl, -N (C₁₋₆-alkyl)₂, -NHC₆₋₁₄aryl, -N(C₆₋₁₄aryl)₂, -N(C₁₋₆alkyl) (C₆₋₁₄aryl), -NHCOR⁶-NO₂, -CN, -F, -Cl, -Br, -I, -O-C₁₋₆ alkyl, -O-C₆₋₁₄-aryl, -O(CO)R⁶, -S-C₁₋₆-alkyl, -S-C₆₋₁₄aryl, -SOR⁶, -SO₃H, -SO₂R⁶, -OSO₂C₁₋₆alkyl, -OSO₂C₆₋₁₄ aryl, -(CS)R⁶, -COOH, -(CO)R⁶ mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycles having 3-14 ring members, mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycles having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S, wherein the C₆₋₁₄aryl groups and the carbocyclic and heterocyclic substituents can optionally be mono- or polysubstituted by R⁴,

R⁵ is

a mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycle having 3-14 ring members or a mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycle having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S, or a carbo- or heterocyclic saturated or mono- or polyunsaturated spirocycle having 3-10 ring members, where heterocyclic systems contain 1-6 heteroatoms, which preferably N, O and S, optionally mono- or polysubstituted by -OH, -SH, -NH₂, -NHC₁₋₆ alkyl, -N(C₁₋₆-alkyl)₂, -NHC₆₋₁₄aryl, -N(C₆₋₁₄aryl)₂,

-N(C₁₋₆alkyl)(C₆₋₁₄aryl), -NHCOR⁶, -NO₂, -CN, -F, -Cl, -Br, -I, -O-C₁₋₅-alkyl, -O-C₆₋₁₄-aryl, -O(CO)R⁶, -S-C₁₋₆-alkyl, -S-C₆₋₁₄aryl, -SOR⁶, -SO₃H, -SO₂R⁶, -OSO₂C₁₋₆alkyl, OSO₂C₆₋₁₄aryl, -(CS)R⁶, -COOH, -(CO)R⁶, mono-, bi or tricyclic saturated or mono- or polyunsaturated carbocycles having 3-14 ring members, mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycles having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S, wherein the C₆₋₁₄aryl groups and the carbocyclic and heterocyclic substituents can optionally be mono- or polysubstituted by R⁴ with the proviso that R⁵ contains at least one substituent selected from -F, -Cl, -Br, -I;

R², R³ are hydrogen or -OH, where at least one of the two substituents must be -OH;

R⁴ is

-H, -OH, -SH, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NHC₆₋₁₄aryl, -N(C₆₋₁₄aryl)₂, -N(C₁₋₆alkyl)(C₆₋₁₄aryl), -NHCOR⁶, -NO₂, -CN, -COOH, -(CO)R⁶, -(CS)R⁶, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -O-C₆₋₁₄-aryl, -O(CO)R⁶, -S-C₁₋₆-alkyl, -S-C₆₋₁₄aryl, -SOR⁶, -SO₂R⁶, -C₁-C₆-alkyl, wherein each aryl or alkyl may be mono- or polysubstituted by -OH, -F, -Cl, -Br, -I;

R⁶ is

-H, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NHC₆₋₁₄aryl, -N(C₆₋₁₄aryl)₂, -N(C₁₋₆alkyl)(C₆₋₁₄aryl), -O-C₁₋₆-alkyl, -O-C₆₋₁₄-aryl, -S-C₁₋₆-alkyl, -S-C₆₋₁₄aryl,

-C₁₋₁₂-alkyl, straight-chain or branched-chain,

-C₂₋₁₂-alkenyl, mono- or polyunsaturated, straight-chain or branched-chain,

mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycles having 3-14 ring members,

mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycles having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S;

A is either a bond, or

$-(CH_2)_m$, $-(CH_2)_m-(CH=CH)_n-(CH_2)_p$, $-(CHOZ)_m$, $-(C=O)-$, $-(C=S)-$, $-(C=N-Z)-$, $-O-$,
 $-S-$, $-NZ-$,

wherein $m, p=0-3$ and $n=0-2$ and

Z is

$-H$, or

$-C_{1-12}$ -alkyl, straight-chain or branched-chain,

$-C_{2-12}$ -alkenyl, mono- or polyunsaturated, straight-chain or branched-chain,

mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycles having 3-14 ring members,

mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycles having 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S;

B is either carbon or sulfur, or $-(S=O)-$;

D is oxygen sulfur, CH_2 or $N-Z$,

where, if B is carbon, D is S or CH_2 ;

E is a bond, or

$-(CH_2)_m$, $-O-$, $-S-$, $-(N-Z)-$, wherein m and Z have the meaning already described above.

2. (previously presented) The method of claim 1 wherein R^5 is selected from monocyclic saturated or mono- or polyunsaturated carbocycles and heterocycles having at least one halogen substituent.

3. (currently amended) The method of claim 2 wherein R^5 is selected from monocyclic aromatic carbocycles and heterocycles having at least one halogen substituent.

4. (previously presented) The method of claim 3 wherein R^5 is a pyridine ring having at least one halogen substituent.

5. (previously presented) The method of claim 3 wherein R^5 is a phenyl ring having at least one halogen substituent.

6. (previously presented) The method of claim 1 wherein R^1 is selected from C_1 - C_{12} alkyl, which is optionally substituted.

7. (previously presented) The method of claim 1 wherein R^1 is selected from monocyclic saturated or mono- or polyunsaturated carbocycles or heterocycles, which are optionally substituted.

8. (previously presented) The method of claim 1 wherein R^2 is OH and R^3 is H.

9. (previously presented) The method of claim 1 wherein A is selected from $-(C=O)-$ and $-(CHOH)-$.

10. (previously presented) The method of claim 1 wherein B is C.

11. (previously presented) The method of claim 1 wherein D is O.

12. (previously presented) The method of claim 1 wherein E is $-(N-H)-$.

13. (previously presented) The method of claim 1 wherein compound (I) is (N-3,5-dichloro-4-pyridinyl)-2-[1-(4-fluorobenzyl)-5-hydroxy-1H-indol-3-yl]-2-oxoacetamide).

14. (canceled)

15. (currently amended) The method of claim 14 wherein the allergic disease is an allergic dermatitis.

16. (canceled)

17. (currently amended) The method of claim 16 wherein the compound is administered to a skin area which is afflicted with the disease after an allergic challenge.

18. (previously presented) The method of claim 17 wherein the compound is administered up to 48 h after the allergic challenge.

19. (canceled)

20. (currently amended) The method of claim 1 ~~19~~ wherein the a further pharmaceutical agent is administered and is a drug that stimulates ~~stimulating~~ cAMP production.

21. (previously presented) The method of claim 20 wherein the further pharmaceutical agent is a corticosteroid.

22. (new) The method of claim 15, wherein the allergic disease is allergic dermatitis.

23. (new) The method of claim 1, further comprising administering a further pharmaceutical agent.